

## MOISTURE TRIGGERED SEALED RELEASE SYSTEM

Cross Reference to Related Application

This application is a Continuation-In-Part of U.S. Patent Application Serial No. 10/122,549, filed April 15, 2002, hereby incorporated in its entirety by reference into this application.

Background of the Invention1. Field of the Invention

The present invention relates to an improved fragrance controlled release system that completely seals the fragrance under ambient conditions, or during the product shelf life such that no fragrance is released until the system is contacted with moisture. More particularly, the invention relates to an improved controlled release system that completely seals menthol under ambient conditions, or during the product shelf life such that no menthol is released until the system is contacted with moisture.

2. Description of the Related Art

Attempts have been made to provide fragrance triggered release systems (in response to moisture) by encapsulating the fragrance in water sensitive materials.

U.S. Patent No. 3,971,852 discloses the use of spray-dried fragrance particles composed of starch derivatives, natural gums (e.g., gum arabic), and polyhydroxy compounds (i.e., mannitol, sorbitol) in cosmetic, personal care, and household products. U.S. Patent No. 4,339,356 also discloses spray-dried fragrance in a water soluble polymeric matrix that emits perfume for a substantial length of time and upon contact with water emits perfume strongly. These patents do not provide any disclosure of selection of selection of types of fragrances for completely sealing the fragrance and improved release upon contact with water.

U.S. Patent Nos. 4,803,195 and 5,508,259 disclose nonaqueous perfuming compositions intended for use in perfumed articles and devices, comprising at least two perfuming elements, wherein each of said perfuming elements has an olfactive character distinct from that of the others, and wherein one of said perfuming elements is in liquid form and the others in water-soluble microencapsulated form. The perfuming composition according to the invention makes it possible to suppress body malodors through topical application thereof on the human body skin and provides advantageous olfactive effects

when used for perfuming soaps and powder detergents. By perfuming element, the invention discloses any active odoriferous ingredient or any perfuming base or mixture of two or more active odoriferous ingredients of current use in perfumery. The matrix material utilized in the above patents comprises:

- 5           a.       a solid film-forming substrate chosen from polyvinyl acetate, polyvinyl alcohol, dextrans, natural or modified starch, vegetable gums, pectins, xanthans, carboxymethylcellulose, methylcellulose, hydroxymethylcellulose and lipoheteropolysaccharides, and
- b.       an emulsifying agent chosen from mono- or diglycerides of fatty acids, esters  
10       derived from the combination of fatty acids with sorbitol or a saccharide, or their alkoxyated derivatives, or an ester of tartaric, citric, ascorbic or lactic acid.

The use of emulsifying agents in these systems increases the solubility of the fragrance in the matrix. The increased solubility of the fragrance in the matrix may affect its permeability through the matrix (C. E. Rogers "Permeation of Gases and Vaours in  
15   Polymers", page 25 in "Polymer Permeability" J. Comyn Ed., incorporated herein as reference):

$$P = D \times S$$

Where:

- P is the fragrance permeability
- 20       D is the diffusion coefficient, and
- S is the solubility

The increased solubility of the fragrance in the matrix as a result of using emulsifying agents may reduce the ability of the matrix to sustain the release of the fragrance from the system during the product shelf life.

- 25       U.S. Patent No. 5,069,231 discloses a controlled release system that provides an initial "burst" of fragrance as well as sustains fragrance release through the use of coating technology. The controlled release system disclosed consists of: (a) a core or a plurality of cores consisting of the same or different releasable functional materials dispersed within a first polymeric substance; (b) coated onto a substantial portion of the surface of the core or  
30       cores, a barrier polymer coating consisting of a second polymeric substance, the second polymeric substance having a permeability to the mass transport of functional material

contained within the core or within one or more of the plurality of cores which is substantially less than the permeability of the first polymeric substance to the mass transport; and (c) coated onto a substantial portion of the outer surface of the barrier polymer coating, a functional material-burst coating consisting of one or more of the same or different functional material dispersed within a third polymeric substance

U.S. Patent Nos. 6,045,835 and 6,106,875 disclose a method of encapsulating flavors and fragrances by controlled water transport into microcapsules. A method of encapsulating an amphiphilic volatile flavor or fragrance compound into a microcapsule having a hydrogel shell and an oil core. The flavor or fragrance compound in a fluid is transported into and solubilized in the core by partition coefficient equilibrium using water in the capsule wall to transport the compound into the core. The fragrances or the flavors are released in response to pressure upon capsule fracture

U.S. Patent No. 6,235,274 discloses microparticles which controllably release olfactorily active substances using extrusion technology. Described are flavor composition, flavor component, perfume composition and perfume component-containing microparticles which are particulate matrices composed of: (a) an olfactorily active component (e.g., perfume component); (b) silica; and (c) a saccharide composition which is a mixture of mannitol and maltose. The microparticles are useful in augmenting, enhancing and/or imparting aroma and/or taste (over relatively long periods of time in a controllably releasable manner) to perfume compositions, perfumed articles (e.g., deodorancy and antiperspirant sticks), foodstuffs, chewing gums, beverages and the like. Also described is a process for preparing the above-mentioned microparticles using, in sequence, (1) adsorption of the olfactorily active material onto silica followed by (2) a blending/extrusion step followed by (3) at least one particularization step.

The prior art of which applicant is aware does not set forth a controlled release system comprising selected fragrances or fragrance ingredients that completely seals the fragrances such that no odor is released until contact with moisture and provides a high impact fragrance burst in response to moisture based on the fragrance ingredient's chemical and physical properties.

### Summary of the Invention

The present invention addresses the ongoing need for fragrance controlled release systems for consumer and cosmetic products that convey to the consumer the product performance by releasing a high intensity odor signal. No engineered coordination of the utilization of the variables concerned has been shown in the prior art whereby, the fragrance ingredients that are utilized for encapsulation in moisture triggered release systems actually have the ability to provide high impact fragrance signal or burst in response to moisture (increase the fraction of the fragrance or fragrance ingredients in the headspace of the system proximate environment after the system has been exposed to moisture). Taking into account that the perception of some fragrance ingredients decreases when exposed to moisture, the careful selection of aroma chemicals to be encapsulated in moisture triggered release systems is critical to effectively deliver a high impact perceivable signal or fragrance burst.

The present invention meets the aforementioned needs in that it has been surprisingly discovered that in order for the moisture triggered release systems to provide high impact fragrance burst it is essential to formulate the fragrance that is encapsulated with fragrance ingredients that have high water solubility and/or high volatility. Suitable fragrance ingredients for encapsulation in the controlled release system of the present invention are naturally, or synthetically, derived fragrance ingredients which have high water solubility as determined by a calculated  $\log_{10}P$ , or  $ClogP \leq$  about 4.0,  $P$  being the n-octanol-water partition coefficient of the fragrance ingredient. The fragrance ingredients can also have high volatility being determined by a boiling point of  $\leq$  about 300° C. The fragrance ingredients solubility in water was found to have a more pronounced effect on their ability to provide a high impact fragrance burst, than the ingredients volatility. A preferred fragrance ingredient is menthol.

The term "fragrance burst" as used herein refers to release of fragrance resulting from release of fragrance ingredients from a moisture-sensitive matrix material upon exposure to water. The term "high impact fragrance burst" as used herein refers to having a high intensity of fragrance immediately released upon exposure of the delivery system to moisture. The "high impact fragrance burst" provides a level of fragrance ingredients in the headspace of the system proximate environment which is much higher than if the encapsulated fragrance comprised fragrance ingredients that have low water solubility and

low volatility (ClogP > 4.0 and boiling point > 300° C). It is believed that fragrance ingredients that have high water solubility and high volatility are more likely to partition or becomes associated with the water through electrostatic interactions and are carried by water vapor into the headspace.

5           The invention also provides a moisture activated controlled release system intended for use in consumer and cosmetic products, such as anhydrous products, that prevent the release of the fragrance during the product shelf life and release the fragrance upon need in response to moisture to provide a high impact fragrance burst.

10           The invention also provides a free-flowing powder comprising micro-spheres composed of water sensitive materials that encapsulate fragrance ingredients that have the desired chemical and physical properties and other active ingredients characterized by:

- (i) protection of the fragrance and the active agents during storage, until needed;
- (ii) moisture triggered release of the active agents upon need in response to moisture; and
- 15       (iii) high impact fragrance burst in response to moisture that "signal" the consumer that the product has been activated.

20           The invention further provides anhydrous consumer and cosmetic products such as deodorant or antiperspirant sticks, deodorant or antiperspirant roll on devices, deodorant or antiperspirant smooth-ons, deodorant or antiperspirant aerosols, body deodorant powders, soap bars, foot spray, hygiene sprays, feminine napkin sprays, undergarment sprays, and the like comprising the controlled release system of the present invention.

          The invention will be more fully described by reference to the following drawings.

#### Brief Description of the Drawings

25           Fig. 1 is a scanning electron microscopy (SEM) with magnification of 5000 times of the micro-spheres of the present invention. The micro-spheres have an average particle size of 10 microns to 20 microns and a smooth surface.

          Fig. 2 is a scanning electron microscopy (SEM) with magnification of 5000 times of the cross section of the micro-spheres of the present invention.

30           Fig. 3 is a graph of the effect of moisture on the content of fragrance ingredients in the headspace.

Fig. 4 is a graph of the effect of moisture on the content of fragrance ingredients in the headspace.

Fig. 5 is a graph of the effect of moisture on the content of fragrance ingredients that have  $\text{ClogP} \leq 4.0$  in the headspace.

5 Detailed Description

The present invention relates to an improved fragrance controlled release system that can be incorporated into a consumer and cosmetic products such as and anhydrous consumer or cosmetic product to provide high intensity odor signal of a high impact fragrance burst in response to moisture. Examples of anhydrous consumer or cosmetic products include  
10 underarm deodorant or antiperspirant sticks, soap bars, body deodorant powders, foot spray, hygiene sprays, feminine napkin sprays, undergarment sprays, and the like. The controlled delivery system of the present invention is a free-flowing powder in the form of micro-spheres composed of water sensitive materials that either dissolves or swells in response to moisture, as shown in Fig. 1 and Fig. 2. The micro-spheres of the present invention have  
15 smooth surface (Fig. 1) and a porous wall (Fig. 2). The term "spheres" is intended to describe solid, substantially spherical particulates. It is appreciated that other particle shapes can be included in the term "sphere" in accordance with the teachings of the present invention.

The fragrance encapsulated in the controlled delivery system of the present invention  
20 is formulated by combining fragrance ingredients that have defined chemical and physical properties, such as high water solubility and high volatility ( $\text{ClogP} \leq 4.0$  and boiling point  $\leq 300^\circ \text{C}$ ) in order to maximize the level of fragrance ingredients in the system proximate environment headspace after the system has been exposed to moisture thereby providing a high odor intensity signal. Suitable encapsulated fragrance ingredients of the present  
25 invention are naturally, or synthetically, derived fragrance ingredients which have high water solubility as determined by a calculated  $\log_{10}P$ , or  $\text{ClogP} \leq 4.0$ ,  $P$  being the n-octanol-water partition coefficient of the fragrance ingredient and high volatility being determined by a boiling point  $\leq 300^\circ \text{C}$ . It will be appreciated that other chemical or physical properties known to those skilled in the art can be used to determine high water solubility and high  
30 volatility.

Additional active ingredients can be added to the controlled release system of the present invention including, but are not limited to: anti-oxidants; free radical scavengers; moisturizers; depigmentation agents; reflectants; humectants; anti-microbial agents, antibacterial agents; allergy inhibitors; anti-acne agents; anti-aging agents; anti-wrinkling agents, antiseptics; analgesics; keratolytic agents; anti-inflammatory agents; fresheners; healing agents; anti infectives; inflammation inhibitors; wound healing promoters; peptides, polypeptides and proteins; deodorants and antiperspirants; skin emollients and skin moisturizers; tanning agents; skin lightening agents; anti-fungal; depilating agents; counterirritants; poison ivy products; poison oak products; burn products; make-up preparations; vitamins; amino acids and their derivatives; herbal extracts; sensory markers; cooling agents; heating agents; skin conditioners; chelating agents; cell turnover enhancers; coloring agents; sunscreens; nourishing agents; moisture absorbers; sebum absorbers and the like; skin penetration enhancers; and other active ingredients. The additional components can be present in an amount of from about 0% to about 20% by weight of the micro-spheres.

#### Selection of fragrance Ingredients for Encapsulation

Fragrances useful for the present invention can be a single aroma chemical, a fragrance accord relatively simple in composition, or can comprise highly sophisticated, complex mixtures of natural and synthetic chemical components, all chosen to provide any desired odor. The encapsulated single aroma chemical, fragrance accord, or full fragrance can be the same or different than a fragrance that is being used as neat oil. The encapsulated fragrance ingredients are preferably selected from the fragrance ingredients comprising the neat fragrance. The encapsulated fragrance is formulated by combining fragrance ingredients that have high water solubility and high volatility. Suitable fragrance ingredients  $\text{ClogP} \leq 4.0$  and/or a boiling point  $\leq 300^\circ \text{C}$  to maximize the level of fragrance ingredients in the system proximate environment headspace after the system has been exposed to moisture to provide a high odor intensity signal. In an embodiment of the present invention, the fragrance ingredient is menthol.

It has been unexpectedly found that the relative amount of fragrance ingredients in the headspace after exposure to water is closely dependent on the ingredient water solubility ( $\text{ClogP}$ ) and/or volatility. The term "headspace" as used herein refers to sampling of a vapor phase above and around the sample. Headspace sampling or headspace analysis ensures that

only volatile materials that are delivered to the vapor phase are analyzed. The solubility in water, or the ability to partition into the aqueous phase, was found to dominate the ability of fragrance ingredients to partition in to the headspace when exposed to water. Fragrance volatility was also observed to affect the ability of fragrance ingredients to partition into the headspace to a lesser extent than the water solubility (ClogP). Figs. 3 and 4 illustrate the effect of moisture on different fragrance ingredients. The fragrance ingredient of limonene, having ClogP of 4.23 and boiling point of 177° C and the fragrance and Benzyl Acetate having ClogP of 1.96 and a boiling point of 215° C were measured in the dry state and in response to moisture, as shown in Fig. 3. The term "au" as used herein refers to arbitrary units. A fragrance ingredient, alpha pinene, having ClogP of 4.6 and boiling point of 156° C and linalool having ClogP of 2.43 and boiling of 198° C were measured in the dry state and in response to moisture, as shown in Fig. 4. Upon exposure to moisture the relative amount of fragrance ingredients that have ClogP > 4.0 in the headspace was observed to decrease as compare to fragrance ingredients that have ClogP < 3.0, although their boiling point was similar in value. Accordingly, when a fragrance is composed primarily of ingredients having a ClogP of about 4.0, or lower and a boiling point of 300° C or lower, the fragrance responds to moisture by increasing its relative amount in the headspace to provide a high impact fragrance burst that can be easily perceived by the consumer. Menthol, another fragrance ingredient, that has ClogP of 3.43 and a boiling point of 216° C was measured in the dry state and in response to moisture. Upon exposure to moisture the relative amount of menthol in the headspace was observed to increase and provided a high impact fragrance burst that can be easily perceived by the consumer.

The boiling points of many perfume ingredients are given in, e.g., "Perfume and Flavor Chemicals (Aroma Chemicals)," S. Arctander, published by the author, 1969, incorporated herein by reference. Other boiling point values can be obtained from different chemistry handbooks and databases, such as the Beilstein Handbook, Lange's Handbook of Chemistry, and the CRC Handbook of Chemistry and Physics. When a boiling point is given only at a different pressure, usually lower pressure than the normal pressure of 760 mm Hg, the boiling point at normal pressure can be approximately estimated by using boiling point-pressure nomographs, such as those given in "The Chemist's Companion," A. J. Gordon and R. A. Ford, John Wiley & Sons Publishers, 1972, pp. 30-36. When applicable, the boiling



point values can also be calculated by computer programs, based on molecular structural data, such as those described in "Computer-Assisted Prediction of Normal Boiling Points of Pyrans and Pyrroles", D. T. Stanton et al, J. Chem. Inf. Comput. Sci., 32 (1992), pp. 306-316, "Computer-Assisted Prediction of Normal Boiling Points of Furans, Tetrahydrofurans, and Thiophenes", D. T. Stanton et al, J. Chem. Inf. Comput. Sci., 31 (1992), pp. 301-310, and references cited therein, and "Predicting Physical Properties from Molecular Structure," R. Murugan et al, Chemtech, June 1994, pp. 17-23. All the above publications are incorporated herein by reference.

The ClogP of many perfume ingredients has been reported; for example, the Pomona92 database, available from Daylight Chemical Information Systems, Inc. (Daylight CIS), Irvine, California, contains many, along with citations to the original literature. However the logP values are most conveniently calculated by the "CLOGP" program, also available from Daylight CIS. This program also lists experimental logP values when they are available in the Pomona92 database. The "calculated logP" (ClogP) is determined by the fragment approach of Hansch and Leo (cf., A. Leo, in Comprehensive Medicinal Chemistry, Volume 4, C. Hansch, P. G. Sammens, J. B. Taylor and C. A. Ramsden, editors, page 295, Pergamon Press, 1990, incorporated by reference herein). The fragment approach is based on the chemical structure of each perfume ingredient and takes into account the numbers and types of atoms, the atom connectivity and the chemical bonding. The ClogP values, which are the most reliable and widely used estimates for this physiochemical property, are preferably used instead of the experimental logP values in the selection of fragrance ingredients which are useful in the present invention.

Fragrance ingredients that are encapsulated in the controlled release system of the present invention to provide high impact fragrance burst in response to moisture are those having a having a ClogP of less than about 4.0, or ClogP of less than about 4.0 and boiling point lower than about 300° C. Table 1 provides nonlimiting examples of fragrance ingredients that can be incorporated in the controlled release system of the present invention to provide a high impact fragrance burst in response to moisture.

Table 1: Examples of bursting fragrance ingredients (ClogP  $\leq$  4.0 and boiling point  $\leq$  300 °C) that can be used in the controlled release system of the present invention

| Aroma Chemical               | Approximate    |       |
|------------------------------|----------------|-------|
|                              | Boiling Points | ClogP |
| Menthol                      | 216            | 3.43  |
| Phenyl ethyl alcohol         | 220            | 1.18  |
| Coumarin                     | 291            | 1.41  |
| Benzaldehyde                 | 179            | 1.48  |
| Hydroxycitronellal           | 241            | 1.54  |
| Vanillin                     | 285            | 1.58  |
| Benzyl acetate               | 215            | 1.96  |
| laevo-Carvone                | 231            | 2.08  |
| Indole                       | 254            | 2.14  |
| beta-Methyl naphthyl ketone  | 300            | 2.28  |
| Eugenol                      | 253            | 2.31  |
| Linalool                     | 198            | 2.43  |
| iso-Eugenol                  | 266            | 2.55  |
| alpha-Terpineol              | 219            | 2.57  |
| Methyl cinnamate             | 263            | 2.62  |
| Geraniol                     | 230            | 2.65  |
| Nerol                        | 227            | 2.65  |
| cis-Jasmone                  | 248            | 2.71  |
| Geranyl acetate              | 245            | 3.72  |
| delta-Nonalactone            | 280            | 2.76  |
| Methyl-N-methyl anthranilate | 256            | 2.79  |
| Dihydro myrcenol             | 208            | 3.03  |
| Benzophenone                 | 306            | 3.12  |
| alpha-Citronellol            | 225            | 3.19  |
| Yara-yara                    | 274            | 3.24  |
| 2-Methoxy naphthalene        | 274            | 3.24  |

| Aroma Chemical               | Approximate    |       |
|------------------------------|----------------|-------|
|                              | Boiling Points | ClogP |
| Phenyl hexanol               | 258            | 3.30  |
| Amyl benzoate                | 262            | 3.42  |
| Phenyl heptanol              | 261            | 3.48  |
| iso-Bornyl acetate           | 227            | 3.49  |
| Linalyl acetate              | 220            | 3.50  |
| delta-Undecalactone          | 290            | 3.83  |
| Lilial                       | 258            | 3.86  |
| Allyl cyclohexane propionate | 267            | 3.93  |

and mixtures thereof.

Fig. 5 illustrates the ability of fragrance ingredients having chemical and physical properties of  $\text{ClogP} \leq 4.0$  and boiling point  $\leq 300^\circ \text{C}$  to provide a burst in response to moisture.

In alternate embodiments of compositions, some fragrance ingredients that have  $\text{ClogP} > \text{about } 4.0$  and boiling point  $> \text{about } 300^\circ \text{C}$  can be used in small amounts, such as to improve product odor. In order to ensure high odor intensity upon exposure of the system to moisture, the fragrance compositions of the present invention contain less than about 30 weight percent of nonbursting fragrance ingredients (fragrance ingredients that have  $\text{ClogP} > \text{about } 4.0$  and boiling point  $> \text{about } 300^\circ \text{C}$ ), preferably less than about 25 weight percent of nonbursting fragrance ingredients, more preferably, less than about 20 weight percent of nonbursting fragrance ingredients, and even more preferably, less than about 15 weight percent of nonbursting fragrance ingredients.

#### Water Sensitive Matrix Materials and Process of Micro-Sphere Formation

Moisture-sensitive materials for forming the micro-spheres of the present invention comprise of water soluble and water dispersible synthetic and natural polymers and copolymers, such as polyvinyl alcohol, starch derivatives, polysaccharides, hydrocolloids, natural gums, proteins, and mixtures thereof. The moisture-sensitive materials can be activated by water or body fluids. The term "body fluids", as used herein, includes eccrine

sweat, apocrine sweat, sebum, build up of sensible moisture from transepidermal water loss, vaginal discharge, urine, and mixtures thereof.

Examples of other synthetic water sensitive polymers which are useful for forming the moisture sensitive of the present invention include polyvinyl pyrrolidone, water soluble  
5 celluloses, ethylene maleic anhydride copolymer, methylvinyl ether maleic anhydride copolymer, acrylic acid copolymers, anionic polymers of methacrylic acid and methacrylate, cationic polymers with dimethyl-aminoethyl ammonium functional groups, polyethylene oxides, water soluble polyamide or polyester.

Examples of water soluble hydroxyalkyl and carboxyalkyl celluloses include  
10 hydroxyethyl and carboxymethyl cellulose, hydroxyethyl and carboxyethyl cellulose, hydroxymethyl and carboxymethyl cellulose, hydroxypropyl carboxymethyl cellulose, hydroxypropyl methyl carboxyethyl cellulose, hydroxypropyl carboxypropyl cellulose, hydroxybutyl carboxymethyl cellulose, and the like. Also useful are alkali metal salts of these carboxyalkyl celluloses, particularly and preferably the sodium and potassium  
15 derivatives.

The polyvinyl alcohol useful in the practice of the invention is partially and fully hydrolyzed polyvinyl acetate, termed "polyvinyl alcohol" with polyvinyl acetate as hydrolyzed to an extent also termed degree of hydrolysis, of from about 75% up to about 99%. Such materials are prepared by means of any of Examples I-XIV of US Patent  
20 No. 5,051,222 issued on September 24, 1991, the specification for which is incorporated by reference herein.

The dissolution rate of polyvinyl alcohol in water is significantly slower than that of maltodextrins and polysaccharide. The dissolution rate of polyvinyl alcohol in water is strongly affected by the polymer degree of hydrolysis and molecular weight. Accordingly,  
25 polyvinyl alcohol can be utilized in the present invention to delay the response of the system to moisture (affect the kinetic of response to moisture), as well as be combined with materials having a faster dissolution rate in order to provide multiple bursts in response to moisture. Polyvinyl alcohol useful for practice of the present invention is Mowiol® 3-83, having a molecular weight of about 14,000 Da and degree of hydrolysis of about 83%, Mowiol® 3-98  
30 and a fully hydrolyzed (98%) polyvinyl alcohol having a molecular weight of 16,000 Da commercially available from Gehring-Montgomery, Inc. of Warminster Pennsylvania.

Other suitable polyvinyl alcohols are: AIRVOL<sup>®</sup> 205, having a molecular weight of about 15,000-27,000 Da and degree of hydrolysis of about 88%, and VINEX<sup>®</sup> 1025, having molecular weight of 15,000-27,000 Da degree of hydrolysis of about 99% and commercially available from Air Products & Chemicals, Inc. of Allentown, Pennsylvania; ELVANOL<sup>®</sup> 51-05, having a molecular weight of about 22,000-26,000 Da and degree of hydrolysis of about 89% and commercially available from the DuPont Company, Polymer Products Department, Wilmington, Delaware; ALCOTEX<sup>®</sup> 78 having a degree of hydrolysis of about 76% to about 79%, ALCOTEX<sup>®</sup> F88/4 having a degree of hydrolysis of about 86% to about 88% and commercially available from the Harlow Chemical Co. Ltd. Of Templefields, Harlow, Essex, England CM20 2BH; and GOHSENOL<sup>®</sup> GL-03 and GOHSENOL<sup>®</sup> KA-20 commercially available from Nippon Gohsei K.K., The Nippon Synthetic Chemical Industry Co., Ltd., of No. 9-6, Nozaki Cho, Kita-Ku, Osaka, 530 Japan.

Suitable polysaccharides are polysaccharides of the nonsweet, coloidally-soluble types, such as natural gums, for example, gum arabic, starch derivates, dextrinized and hydrolyzed starches, and the like. A suitable polysaccharide is a water dispersible, modified starch commercially available as Capul<sup>®</sup>, N-Lok<sup>®</sup>, Hi-Cap<sup>™</sup> 100 or Hi-Cap<sup>™</sup> 200 commercially available from the National Starch and Chemical Company of Bridgewater, New Jersey; Pure-Cote<sup>™</sup>, commercially available from the Grain Processing Corporation of Muscatine, Iowa. In the preferred embodiment, the natural gum is a gum arabic, commercially available from TIC Gums Inc. Belcamp, Midland. Suitable hydrocolloids are xanthan, maltodextrin, galactomanan or tragacanth, preferably maltodextrins such as Maltrin<sup>™</sup> M100, and Maltrin<sup>™</sup> M150, commercially available from the Grain Processing Corporation of Muscatine, Iowa.

The amount of fragrance based in moisture sensitive micro-sphere can be in the range of from about 1% up to about 60% by weight of the moisture sensitive micro-sphere.

The micro-spheres of the present invention can be prepared by the steps of (1) forming an aqueous phase of the moisture sensitive materials (either a single material or mixture of several materials); (2) emulsifying the fragrance, fragrance accord, or fragrance ingredient and other active agents in the aqueous phase; and (3) removing moisture to create free-flowing powder. For example, moisture can be removed by spray drying droplets of emulsion. Spray drying is well known in the art and been used commercially in many

applications, including foods where the core material is a flavoring oil and cosmetics where the core material is a fragrance oil, as described in Cf. Balassa, "Microencapsulation in the Food Industry", CRC Critical Review Journal in Food Technology, July 1971, pp 245-265; Barreto, "Spray Dried Perfumes for Specialties, Soap and Chemical Specialties",  
5 December 1966; Maleeny, Spray Dried Perfumes, Soap and San Chem, Jan. 1958, pp. 135 et seq.; Flinn and Nack, "Advances in Microencapsulation Techniques", Batelle Technical Review, Vo. 16, No. 2, pp. 2-8 (1967); US patent Nos. 5,525,367; and 5,417,153 which are incorporated herein as references.

In one embodiment, micro-spheres that have the ability to provide multiple fragrance  
10 bursts in response to moisture are formed by mixing the fragrance and a selected active agent with polyvinyl alcohol, or compositions of polyvinyl alcohol and polysaccharides, under conditions sufficient to encapsulate the fragrance. Preferably mixing a selected fragrance and active agent with the polyvinyl alcohol, or compositions of polyvinyl alcohol and polysaccharides, until the emulsion is formed and then spray drying the emulsion to thereby  
15 form an encapsulated micro-sphere. In the preferred embodiment, the moisture sensitive matrix is formed of a polyvinyl alcohol material at a level of from about 1% to about 80%, preferably from about 1% to about 70% by weight of the matrix material with the balance being the amount by weight of active agents and an optimal amount of polysaccharides. In an alternate embodiment, polyvinyl alcohol is present in the matrix material in an amount of  
20 about 1% to about 80% by weight and polysaccharides are present in the matrix material in an amount of about 1% to about 80% by weight. In the preferred embodiment, the fragrance ingredients are generally present at a level from about 0.01% to about 80% preferably from about 1% to about 50% by weight of the encapsulated active agent with the balance being the polyvinyl alcohol or polyvinyl alcohol and polysaccharides. Optionally other conventional  
25 ingredients known in the art such as preservatives can be used in accordance with the teachings of the present invention. The micro-spheres of the present invention preferably have size of from about 0.5 micron to about 300 microns, more preferably from about 1 micron to about 200 microns, most preferably from about 2 microns to about 30 microns. The present invention preferably has minimal active agents on the surface of the spheres,  
30 preferably less than 1%.

The invention can be further illustrated by the following examples thereof, although it will be understood that these examples are included merely for purposes of illustration and are not intended to limit the scope of the invention unless otherwise specifically indicated. All percentages, ratios, and parts herein, in the Specification, Examples, and Claims, are by

5 weight and are approximations unless otherwise stated.

## THE NEAT FRAGRANCE OIL

EXAMPLE 1

5 The composition of the fragrance that was used as neat oil is detailed to the following table. The neat fragrance oil is composed of a wide range of fragrance ingredients including both bursting and nonbursting ingredients.

| Fragrance Composition | Component (%Wt.) | ClogP | Boiling Point (°C) |
|-----------------------|------------------|-------|--------------------|
| Alpha Pinene          | 15.0             | 4.6   | 198                |
| Geraniol              | 5.0              | 2.65  | 230                |
| Dihydro Myrcenol      | 5.0              | 3.03  | 208                |
| Linalool              | 5.0              | 2.43  | 198                |
| Lemonene              | 20.0             | 4.23  | 177                |
| alpha-Citronellol     | 5.0              | 3.19  | 225                |
| Phenyl Ethyl Alcohol  | 5.0              | 2.57  | 219                |
| Linalyl benzoate      | 10.0             | 5.233 | 263                |
| Galaxolide            | 30.0             | 5.482 | +250               |



## CREATION OF A FRAGRANCE ACCORD FOR ENCAPSULATION

### EXAMPLE 2

The fragrance ingredients for encapsulation have  $\text{ClogP} \leq 4.0$  and boiling point  $\leq$   
 5 300° C and are selected from the composition of the neat fragrance.

| Fragrance Composition | Component (%Wt.) | ClogP | Boiling Point (°C) |
|-----------------------|------------------|-------|--------------------|
| Geraniol              | 30.0             | 2.65  | 230                |
| Dihydro Myrcenol      | 20.0             | 3.03  | 208                |
| Linalool              | 15.0             | 2.43  | 198                |
| alpha-Citronellol     | 5.0              | 3.19  | 225                |
| Phenyl Ethyl Alcohol  | 30.0             | 2.57  | 219                |

## PREPARATION OF DUAL FRAGRANCE CONTROLLED RELEASE SYSTEM

### 10 EXAMPLE 3

The following procedure is used for the preparation of a controlled release system that provides high impact fragrance burst in response to moisture. The micro-sphere water sensitive matrix is a mixture of Hi-Cap™ 100 (commercially available from the National Starch and Chemical Company of Bridgewater, New Jersey), modified starch, Pure-Cote™,  
 15 commercially available from the Grain Processing Corporation of Muscatine, Iowa, and maltodextrin, Maltrin™ M100, commercially available from the Grain Processing Corporation of Muscatine, Iowa.

400 grams of the fragrance accord of Example 2 are emulsified with 240 grams of Hi-Cap™ 100 (commercially available from the National Starch and Chemical Company of  
 20 Bridgewater, New Jersey), 100 grams of Pure-Cote™ (commercially available from the Grain Processing Corporation of Muscatine, Iowa), 60 grams of Maltrin™ M100 (commercially available from the Grain Processing Corporation of Muscatine, Iowa), and 1200 grams of water. The resulting emulsion is spray dried with a Bowen Lab Model Drier (at Spray-Tek of Middlesex, New Jersey) utilizing 250 c.f.m of air with an inlet temperature

of 380 °F, and outlet temperature of 225 °F and a wheel speed of 45,000 r.p.m to produce a free flowing, dry powder, consisting of 50% encapsulated fragrance.

#### EXAMPLE 4

400 grams of the fragrance accord of Example 1 are emulsified with 240 grams of Hi-Cap™ 100 (commercially available from the National Starch and Chemical Company of Bridgewater, New Jersey), 100 grams of Pure-Cote™ (commercially available from the Grain Processing Corporation of Muscatine, Iowa), 60 grams of Maltrin™ M100 (commercially available from the Grain Processing Corporation of Muscatine, Iowa), and 1200 grams of water. The resulting emulsion is spray dried with a Bowen Lab Model Drier (at Spray-Tek of Middlesex, New Jersey) utilizing 250 c.f.m of air with an inlet temperature of 380 °F, and outlet temperature of 225 °F and a wheel speed of 45,000 r.p.m to produce a free flowing, dry powder, consisting of 50% encapsulated fragrance.

#### EXAMPLE 5

The following procedure is used for the preparation of a controlled release system that provides high impact fragrance burst in response to moisture. The micro-sphere water sensitive matrix is polyvinyl alcohol a having a molecular weight of about 14,000 Da and degree of hydrolysis of about 83% (Mowiol® 3-83, trademark and commercially available from Gehring–Montgomery, Inc. of Warminster Pennsylvania).

200 grams of the fragrance accord of Example 2 are emulsified with 400 grams polyvinyl alcohol a having a molecular weight of about 14,000 Da and degree of hydrolysis of about 83% (Mowiol® 3-83, trademark and commercially available from Gehring–Montgomery, Inc. of Warminster Pennsylvania) and 1400 grams of water. The resulting emulsion is spray dried with a Bowen Lab Model Drier (at Spray-Tek of Middlesex, New Jersey) utilizing 250 c.f.m of air with an inlet temperature of 380 °F, and outlet temperature of 225 °F and a wheel speed of 45,000 r.p.m to produce a free flowing, dry powder, consisting of 30% encapsulated fragrance.

#### EXAMPLE 6

200 grams of the fragrance accord of Example 1 are emulsified with 400 grams polyvinyl alcohol a having a molecular weight of about 14,000 Da and degree of hydrolysis of about 83% (Mowiol® 3-83, trademark and commercially available from Gehring–Montgomery, Inc. of Warminster Pennsylvania) and 1400 grams of water. The resulting

emulsion is spray dried with a Bowen Lab Model Drier (at Spray-Tek of Middlesex, New Jersey) utilizing 250 c.f.m of air with an inlet temperature of 380 °F, and outlet temperature of 225 °F and a wheel speed of 45,000 r.p.m to produce a free flowing, dry powder, consisting of 30% encapsulated fragrance.

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#### EXAMPLE 7

The following procedure is used for the preparation of a controlled release system that provides high impact fragrance burst in response to moisture. The micro-sphere water sensitive matrix is a mixture of polyvinyl alcohol a having a molecular weight of about 14,000 Da and degree of hydrolysis of about 83% (Mowiol® 3-83, trademark and commercially available from Gehring–Montgomery, Inc. of Warminster Pennsylvania) and Hi-Cap™ 200 (commercially available from the National Starch and Chemical Company of Bridgewater, New Jersey).

200 grams of the fragrance accord of Example 2 are emulsified with 20 grams polyvinyl alcohol a having a molecular weight of about 14,000 Da and degree of hydrolysis of about 83% (Mowiol® 3-83, trademark and commercially available from Gehring–Montgomery, Inc. of Warminster Pennsylvania), 360 grams of Hi-Cap™ 200 (commercially available from the National Starch and Chemical Company of Bridgewater, New Jersey), and 1200 grams of water. The resulting emulsion is spray dried with a Bowen Lab Model Drier (at Spray-Tek of Middlesex, New Jersey) utilizing 250 c.f.m of air with an inlet temperature of 380 °F, and outlet temperature of 225 °F and a wheel speed of 45,000 r.p.m to produce a free flowing, dry powder, consisting of 50% encapsulated fragrance.

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#### EXAMPLE 8

200 grams of menthol is emulsified with 800 grams polyvinyl alcohol a having a molecular weight of about 14,000 Da and degree of hydrolysis of about 83% (Mowiol® 3-83, trademark and commercially available from Gehring–Montgomery, Inc. of Warminster Pennsylvania) and 1500 grams of water. The resulting emulsion is spray dried with a Bowen Lab Model Drier (at Spray-Tek of Middlesex, New Jersey) utilizing 250 c.f.m of air with an inlet temperature of 380 °F, and outlet temperature of 225 °F and a wheel speed of 45,000 r.p.m to produce a free flowing, dry powder, consisting of 20% encapsulated menthol.

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## INCORPORATION OF THE CONTROLLED RELEASE SYSTEM IN UNDERARM PRODUCTS

### EXAMPLE 9

5 The performance of an antiperspirant product comprising the fragrance controlled release system of Examples 3 to Example 8 (i.e., the ability to yield a high impact multiple fragrance "burst" upon perspiration) was evaluated using 0.5% neat fragrance oil of Example 1 and 1.5% encapsulated fragrance (encapsulated nonbursting fragrance (Example 1) or encapsulation of bursting fragrance (Example 2)). The un-fragranced antiperspirant base was a commercial Degree® unscented antiperspirant stick available from  
10 Helene Curtis Company of Chicago, Illinois that is fragrance free.

The control samples were prepared by weighting into a jar the appropriate amount of the neat fragrance of example 1 or the encapsulated nonbursting fragrances and the appropriate amount of the unscented Degree® base and the resulting mixture was melted at 80° C and mixed for about one minute. The melt was poured into an underarm container and  
15 allowed to cool for an hour. The antiperspirant stick comprising the controlled release system of the present invention is prepared by weighting the appropriate amount of the neat fragrance of Example 1 the appropriate amount of the encapsulated fragrance powder and the unscented Degree® base melt into a jar. The melt was poured into an underarm container and allowed to cool for an hour.

20 The antiperspirant samples were applied on the forearm. The ability of the products to provide multiple fragrance burst was evaluated one hour after application of the product by misting the area with water. The products comprising the controlled release system of the present invention were found to provide high impact fragrance burst upon wetting the area whereas the control sample comprising the neat oil did not. The forearm areas treated with  
25 the antiperspirant comprising the neat fragrance (control) and that treated with the antiperspirant comprising the encapsulated fragrance were evaluated for their ability to provide high impact burst again after 6 hours (provide multiple fragrance burst). The ability of the systems to provide high impact fragrance burst was determined by olfactive evaluation. Odor perception is, by its nature, a very subjective determination. According to  
30 the procedure, the samples to be tested are provided to a panel of six odor specialists who independently rank the odor intensity on a scale of 1 (least) to 10 (most) for odor and

intensity. Samples yielding an odor ranking below about 3.0 possess an odor which would hardly be noticed by the general public. The odor evaluation results were as follow:

|                                    | Response to Moisture   |                         |
|------------------------------------|------------------------|-------------------------|
|                                    | Evaluated after 1 Hour | Evaluated after 6 Hours |
| Neat Fragrance Oil (Example 1)     | 4                      | 2                       |
| Encapsulated Fragrance (Example 3) | 7                      | 5                       |
| Encapsulated Fragrance (Example 4) | 4                      | 3                       |
| Encapsulated Fragrance (Example 5) | 6                      | 6                       |
| Encapsulated Fragrance (Example 6) | 3                      | 4                       |
| Encapsulated Fragrance (Example 7) | 8                      | 5                       |
| Encapsulated Fragrance (Example 8) | 4                      | 4                       |

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These results clearly show that only the forearm treated with the encapsulated bursting fragrance of Example 2 comprising fragrance ingredients with  $\text{ClogP} \leq 4.0$  and boiling point  $\leq 300^\circ \text{C}$  provide multiple high impact fragrance burst that can be easily perceived. The forearm treated with the neat fragrance or the encapsulated nonbursting fragrance of Example 1 had very low intensity signal in response to moisture.

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#### INCORPORATION OF THE CONTROLLED RELEASE SYSTEM IN BODY DEODORANT POWDER

##### EXAMPLE 10

The performance of a deodorant body powder comprising the fragrance controlled release system of Example 3-8 (i.e., the ability to yield a high impact fragrance "burst" upon perspiration) was evaluated and compared to the performance of the same product comprising the neat fragrance, at the same fragrance level. The un-fragranced body deodorant powder base was unscented Pure-Dent® commercially available from Grain Processing Corporation of Muscatine, Iowa.

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The body deodorant powders were prepared at a 1% effective fragrance concentration using the fragrance described in Example 1 and Example 2. The control samples was prepared by weighting into a jar appropriate amounts of the neat fragrance or encapsulated

fragrance of Example 4, Example 6, and Example 8 appropriate amounts of the unscented Pure-Dent® powder followed by mixing the powder well. The body deodorant powder comprising the controlled release system of the present invention is prepared by weighting appropriate amounts of the unscented Pure-Dent® powder into a jar following by adding  
 5 appropriate amounts of the powder of Example 3, Example 5, and Example 7, following by mixing the powders well.

The body deodorant powders samples were applied on the forearm. The ability of the products to provide multiple fragrance burst was evaluated one hour and after 6 hours following application of the product by misting the area with water. The product comprising  
 10 the controlled release system of the present invention was found to provide high impact fragrance burst upon wetting the forearm area with water whereas the control samples comprising the neat oil or the encapsulated nonbursting fragrance did not.

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|                                    | Response to Moisture   |                         |
|------------------------------------|------------------------|-------------------------|
|                                    | Evaluated after 1 Hour | Evaluated after 6 Hours |
| Neat Fragrance Oil (Example 1)     | 4                      | 3                       |
| Encapsulated Fragrance (Example 3) | 7                      | 5                       |
| Encapsulated Fragrance (Example 4) | 4                      | 2                       |
| Encapsulated Fragrance (Example 5) | 6                      | 6                       |
| Encapsulated Fragrance (Example 6) | 3                      | 2                       |
| Encapsulated Fragrance (Example 7) | 8                      | 5                       |
| Encapsulated Fragrance (Example 8) | 4                      | 3                       |

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